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     1
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NEWS
     2
                New IPC8 SEARCH, DISPLAY, and SELECT fields in USPATFULL/
NEWS
     3 DEC 23
                USPAT2
        JAN 13
                IPC 8 searching in IFIPAT, IFIUDB, and IFICDB
NEWS
                New IPC 8 SEARCH, DISPLAY, and SELECT enhancements added to
NEWS 5
        JAN 13
                INPADOC
                Pre-1988 INPI data added to MARPAT
NEWS 6
        JAN 17
                IPC 8 in the WPI family of databases including WPIFV
NEWS 7
        JAN 17
                Saved answer limit increased
NEWS 8
        JAN 30
                STN AnaVist, Version 1.1, lets you share your STN AnaVist
NEWS 9
        FEB 21
                visualization results
                The IPC thesaurus added to additional patent databases on STN
NEWS 10 FEB 22
                Updates in EPFULL; IPC 8 enhancements added
NEWS 11
        FEB 22
                New STN AnaVist pricing effective March 1, 2006
NEWS 12 FEB 27
                MEDLINE/LMEDLINE reload improves functionality
NEWS 13 FEB 28
                TOXCENTER reloaded with enhancements
NEWS 14 FEB 28
                REGISTRY/ZREGISTRY enhanced with more experimental spectral
NEWS 15 FEB 28
                property data
                INSPEC reloaded and enhanced
NEWS 16 MAR 01
                Updates in PATDPA; addition of IPC 8 data without attributes
NEWS 17 MAR 03
                X.25 communication option no longer available after June 2006
NEWS 18 MAR 08
                EMBASE is now updated on a daily basis
NEWS 19 MAR 22
                New IPC 8 fields and IPC thesaurus added to PATDPAFULL
NEWS 20 APR 03
                Bibliographic data updates resume; new IPC 8 fields and IPC
NEWS 21 APR 03
                 thesaurus added in PCTFULL
                 STN AnaVist $500 visualization usage credit offered
NEWS 22 APR 04
                LINSPEC, learning database for INSPEC, reloaded and enhanced
NEWS 23 APR 12
                 Improved structure highlighting in FQHIT and QHIT display
NEWS 24 APR 12
                 in MARPAT
                 Derwent World Patents Index to be reloaded and enhanced during
NEWS 25
        APR 12
                 second quarter; strategies may be affected
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NEWS EXPRESS FEBRUARY 15 CURRENT VERSION FOR WINDOWS IS V8.01a,
CURRENT MACINTOSH VERSION IS V6.0c(ENG) AND V6.0Jc(JP),
AND CURRENT DISCOVER FILE IS DATED 19 DECEMBER 2005.
V8.0 AND V8.01 USERS CAN OBTAIN THE UPGRADE TO V8.01a AT http://download.cas.org/express/v8.0-Discover/

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=> FIL CAPLUS
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SINCE FILE TOTAL ENTRY SESSION 0.46 0.67

FULL ESTIMATED COST

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FILE COVERS 1907 - 27 Apr 2006 VOL 144 ISS 18 FILE LAST UPDATED: 26 Apr 2006 (20060426/ED)

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     ANSWER 1 OF 1 CAPLUS COPYRIGHT 2006 ACS on STN
L1
     2004:120553 CAPLUS
ΑN
DN
     140:184417
     Bicyclic thioureas as lubricating oil antiwear, anticorrosion, and
ΤI
     antioxidant additives to replace zinc dialkyl dithiophosphates
     Mukkamala, Ravindranath
IN
PA
SO
     U.S. Pat. Appl. Publ., 4 pp.
     CODEN: USXXCO
DT
     Patent
LA
     English
FAN.CNT 1
                                          APPLICATION NO.
                        KIND
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                                                                 DATE
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                                          US 2003-636111
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L2
=> DIS L2 1 IBIB IABS
THE ESTIMATED COST FOR THIS REQUEST IS 2.74 U.S. DOLLARS
DO YOU WANT TO CONTINUE WITH THIS REQUEST? (Y) /N:Y
     ANSWER 1 OF 34 CAPLUS COPYRIGHT 2006 ACS on STN
L2
                         2004:120553 CAPLUS
ACCESSION NUMBER:
                         140:184417
DOCUMENT NUMBER:
                         Bicyclic thioureas as lubricating oil antiwear,
TITLE:
                         anticorrosion, and antioxidant additives to replace
                         zinc dialkyl dithiophosphates
                         Mukkamala, Ravindranath
INVENTOR(S):
PATENT ASSIGNEE(S):
                         USA
                         U.S. Pat. Appl. Publ., 4 pp.
SOURCE:
                         CODEN: USXXCO
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DOCUMENT TYPE:

Patent

LANGUAGE:

English

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 2004029743	A1	20040212	US 2003-636111	20030807
EP 1394241	A1	20040303	EP 2003-254689	20030728
R: AT, BE, CH,	DE, DK	, ES, FR, GE	B, GR, IT, LI, LU,	NL, SE, MC, PT,
IE, SI, LT,	LV, FI	, RO, MK, CY	(, AL, TR, BG, CZ,	EE, HU, SK
JP 2004162027	A2	20040610	JP 2003-287826	20030806
PRIORITY APPLN. INFO .:			US 2002-401540P	P 20020807

OTHER SOURCE(S):

MARPAT 140:184417

GRAPHIC IMAGE:

ABSTRACT:

Cyclic and bicyclic thiourea-based lubricating oil antiwear, corrosion inhibitor, and antioxidant additives, as suitable replacements for zinc dialkyl dithiophosphates, are of general structure I, in which W = O or S; R1 and R2 are H, alkyl, alkenyl, aryl, or aralkyl; R3 and R4 are H, alkyl, alkenyl, aryl, or aralkyl, or R3 and R4 combine with ring carbon atoms to which they are attached to form a five-, six-, or seven-membered heterocyclic ring. At least one of R1-4 is C6-22-alkyl. In addition, the additives can be of general structure II, in which R1 and R2 are as described above, and R5. and R6 are H, alkyl, alkenyl, aryl, or aralkyl. The additives are present at 0.1-20 weight% concentration

=> DIS L2 2 IBIB IABS

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ANSWER 2 OF 34 CAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER:

2001:107608 CAPLUS

DOCUMENT NUMBER:

134:203693

TITLE:

Biorational design, synthesis, and inhibition of

photosystem II inhibitors

AUTHOR (S):

Liu, Huayin; Sha, Yinlin; Yu, Aiming; Tan, Huifen;

Yang, Huazheng

CORPORATE SOURCE:

State Key Laboratory of Elemento-Organic Chemistry,

Institute of Elemento-Organic Chemistry, Nankai University, Tianjin, 300071, Peop. Rep. China ACS Symposium Series (2001), 774 (Agrochemical

SOURCE: Discovery), 226-235

CODEN: ACSMC8; ISSN: 0097-6156

PUBLISHER: DOCUMENT TYPE: American Chemical Society Journal; General Review

English

LANGUAGE: ABSTRACT:

A review with 19 refs. Mol. modeling of cyanoacrylates(cyanoacrylamides) with D1 protein of Pisum sativum have been presented. Studies show that the binding force includes mainly: H-bond interaction, Van der Waals and π -ring stacking interaction. It was found that SER 268 in D1 protein might be an important binding site. Thus some new cyanoacrylates (cyanoacrylamides) were designed and synthesized. For rapid optimization, combinatorial methods were introduced to synthesize 3-N-substituted (2-thio)hydrouracils library by the acidic cyclization-cleavage of ureas and thioureas from Wang resin. Their Hill inhibition is discussed.

REFERENCE COUNT: 19 THERE ARE 19 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

=> DIS L2 3 IBIB IABS

THE ESTIMATED COST FOR THIS REQUEST IS 2.74 U.S. DOLLARS DO YOU WANT TO CONTINUE WITH THIS REQUEST? (Y)/N:Y

L2 ANSWER 3 OF 34 CAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 1998:250700 CAPLUS

DOCUMENT NUMBER: 128:295059

TITLE: Preparation of pyridyl- and naphthyridylalkoxybenzoyl-

 $\alpha\text{-}(\text{phenylsulfonylamino})\text{-}\beta\text{-}\text{alanine}$ derivatives and analogs for inhibiting osteoclast-mediated bone resorption

INVENTOR(S): Hartman, George D.; Duggan, Mark E.; Hoffman, William

F.; Ihle, Nathan C.

PATENT ASSIGNEE(S): Merck and Co., Inc., USA

SOURCE: U.S., 57 pp., Cont.-in-part of U.S. Ser. No. 250,218,

abandoned.

CODEN: USXXAM

DOCUMENT TYPE: Patent LANGUAGE: English

FAMILY ACC. NUM. COUNT: 2

PATENT INFORMATION:

PATENT NO.	KIND DATE	APPLICATION NO.	DATE
02 0 1 1 1 1 1 1	A 19980421		
WO 9532710	A1 19951207	WO 1995-US5938	19950512
W: AM, AU, BB,	BG, BR, BY, CA,	CN, CZ, EE, FI, GE,	HU, IS, JP, KG,
		MG, MN, MX, NO, NZ,	
SI, SK, TJ,	TM, TT, UA, US,	UZ	
RW: KE, MW, SD,	SZ, UG, AT, BE,	CH, DE, DK, ES, FR,	GB, GR, IE, IT,
LU, MC, NL,	PT, SE, BF, BJ,	CF, CG, CI, CM, GA,	GN, ML, MR, NE,
SN, TD, TG			
US 5929120	A 19990727	US 1998-15982	19980130
PRIORITY APPLN. INFO.:		US 1994-250218	B2 19940527
		WO 1995-US5938	W 19950512
		US 1996-714097	A3 19960926

OTHER SOURCE(S): MARPAT 128:295059

GRAPHIC IMAGE:

ABSTRACT:

Compds. of structure I [X = various amino, amidino, guanidino, and N-heterocyclic groups; Y = alkylene, alkynylene, alkenylene, etc.; B = alkylene with optional amide moiety in chain; R1 = H, alkoxyalkyl, alkoxycarbonylalkyl, (di)(alkyl)aminoalkyl, aralkyl; R6, R7 = H, (di)alkylaminoalkyl, alkoxycarbonylaminoalkyl, alkylsulfonylaminoalkyl, alkylcarbonylaminoalkyl; R12 = OH, alkoxy, dialkylaminocarbonylmethoxy, aryldialkylaminocarbonylmethoxy; with provisos), are described which inhibit osteoclast-mediated bone resorption. Specifically, the compds. are useful for treating mammals suffering from a bone condition caused or mediated by increased bone resorption, who are in need of such therapy. The compds. may be administered in oral dosage forms such as tablets, capsules, e.g. sustained release capsules, powders, granules, and suspensions. Syntheses of approx. 50 compds. in 37 synthetic examples are described. Thus, amidation of Me 4-[2-(4-aminopyridin-6-yl)ethoxy]benzoic acid (preparation given) with (R)-H2NCH2CH(NHSO2Ph)CO2CMe3.HCl (preparation given) using EDC, Nhydroxybenzotriazole (HOBt), and N-methylmorpholine in DMF, followed by deprotection with CF3CO2H gave desired compound II. In EIB and OCFORM assays, prepared compds. I had values ranging 0.5-500 nM and 1-1000 nM, resp.

REFERENCE COUNT: 6 THERE ARE 6 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

=> DIS L2 4 IBIB IABS
THE ESTIMATED COST FOR THIS REQUEST IS 2.74 U.S. DOLLARS
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L2 ANSWER 4 OF 34 CAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER:

1996:181547 CAPLUS

DOCUMENT NUMBER:

124:232066

TITLE:

N-(Guanidinoalkoxybenzoyl)- α -

(phenylsulfonylamino)- β -alanine derivatives and analogs for inhibiting osteoclast-mediated bone

resorption

INVENTOR(S):

Hartman, George D.; Duggan, Mark E.; Ihle, Nathan C.;

Hoffman, William F.

PATENT ASSIGNEE(S):

Merck and Co., Inc., USA PCT Int. Appl., 241 pp.

CODEN: PIXXD2

DOCUMENT TYPE:

Patent

LANGUAGE:

SOURCE:

English

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 9532710	A1	19951207	WO 1995-US5938	19950512

W: AM, AU, BB, BG, BR, BY, CA, CN, CZ, EE, FI, GE, HU, IS, JP, KG, KR, KZ, LK, LR, LT, LV, MD, MG, MN, MX, NO, NZ, PL, RO, RU, SG, SI, SK, TJ, TM, TT, UA, US, UZ RW: KE, MW, SD, SZ, UG, AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG 19951207 CA 1995-2190870 19950512 CA 2190870 AΑ A1 AU 1995-25868 19950512 AU 9525868 19951221 B2 19990204 AU 701776 EP 1995-920409 19950512 A1 19970312 EP 760658 EP 760658 B1 20021113 R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LI, LU, NL, PT, SE T2 JP 1996-500899 19950512 JP 10501222 19980203 E AT 1995-920409 19950512 AT 227567 20021115 **T3** ES 1995-920409 19950512 ES 2186720 20030516 19980421 US 1996-714097 19960926 US 5741796 Α US 1994-250218 A 19940527 PRIORITY APPLN. INFO.: WO 1995-US5938 W 19950512

OTHER SOURCE(S): GRAPHIC IMAGE: MARPAT 124:232066

$$\begin{array}{c|c}
R^1 & O \\
\hline
 & R^{12} \\
\hline
 & R^{6} & R^{7}
\end{array}$$

ABSTRACT:

Compds. of structure I [X = various amino, amidino, guanidino, and N-heterocyclic groups; Y = alkylene, alkynylene, alkenylene, etc.; B = alkylene with optional amide moiety in chain; R1 = H, alkoxyalkyl, alkoxycarbonylalkyl, (di)(alkyl)aminoalkyl, aralkyl; R6, R7 = H, (di)alkylaminoalkyl, alkoxycarbonylaminoalkyl, alkylsulfonylaminoalkyl, alkylcarbonylaminoalkyl; R12 = OH, alkoxy, dialkylaminocarbonylmethoxy, aryldialkylaminocarbonylmethoxy; with a proviso], which inhibit osteoclast-mediated bone resorption. Syntheses of approx. 50 compds. in 37 synthetic examples are described. For example, amidation of 4-(BOC-NHCH2CH2O)C6H4CO2H with (R)-H2NCH2CH(NHSO2Ph)CO2Bu-tert.HCl [preparation given] using BOP reagent and NMM in MeCN, followed by deprotection with CF3CO2H and condensation of the amine with DPFN [3,5-dimethyl-1-pyrazolylformamidine nitrate], gave title compound II. In the EIB and OCFORM assays, I had values ranging 0.5-500 nM and 1-1000 nM, resp.

II

=> DIS L2 5 IBIB IABS
THE ESTIMATED COST FOR THIS REQUEST IS 2.74 U.S. DOLLARS
DO YOU WANT TO CONTINUE WITH THIS REQUEST? (Y)/N:Y

L2 ANSWER 5 OF 34 CAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 1995:605449 CAPLUS

DOCUMENT NUMBER: 123:127424

TITLE: Method for processing a black-and-white silver halide

photographic light-sensitive material.

INVENTOR(S): Fukawa, Junichi; Goto, Kenji; Sampei, Takeshi

PATENT ASSIGNEE(S): Konica Corp., Japan SOURCE: Eur. Pat. Appl., 63 pp.

CODEN: EPXXDW

DOCUMENT TYPE: Patent LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
EP 631179	Al	19941228	EP 1994-303803	19940526
EP 631179	Bl	20000405		
R: DE, FR, GB,	IT			
JP 06347953	A2	19941222	JP 1993-138704	19930610
JP 3350739	B2	20021125		
US 5441847	Α	19950815	US 1994-249455	19940526
PRIORITY APPLN. INFO.:			JP 1993-138704 A	19930610
OTHER SOURCE(S):	MARPAT	123:127424		
GRAPHIC IMAGE:				

ABSTRACT:

A method is described of processing, using an automatic processor, a black-and-white Ag halide photog. light-sensitive material containing a hydrazine compound and a redox compound capable of releasing an inhibitor upon oxidation, wherein a developer is replenished by a developer replenisher at a rate of $\leq 200 \text{ mL/m2}$ of the photog. material, and the developer has a pH of 9.5 to 10.8. The developer further contains Z-SM, I, or II [Z = alkyl, aryl, or heterocyclyl each of which has ≥ 1 group selected from OH, SO3M1, CO2M1, amino, ammonio, or a group containing ≥ 1 of the above group as substituent; M = H, alkali metal, amidino; M1 = H, alkali metal, ammonium; Y21 and Z21, Y31 and Z31 = atoms necessary to form a ring, and thus formed ring contain ≥ 3 N atoms and are substituted by a mercapto group]. The material produces high contrast images without deteriorating the sensitivity and the without producing spots.

=> DIS L2 6 IBIB IABS

THE ESTIMATED COST FOR THIS REQUEST IS 2.74 U.S. DOLLARS DO YOU WANT TO CONTINUE WITH THIS REQUEST? (Y)/N:Y

L2 ANSWER 6 OF 34 CAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 1986:88492 CAPLUS

DOCUMENT NUMBER: 104:88492

TITLE: Heterocycles. 79. N1- and N2-substituted

2-amino-5,6-dihydro-4(1H)-pyrimidinones

AUTHOR(S): Wendelin, Winfried; Riedl, Renate

CORPORATE SOURCE: Inst. Pharm. Chem., Univ. Graz, Graz, A-8010, Austria

SOURCE: Monatshefte fuer Chemie (1985), 116(2), 237-51

CODEN: MOCMB7; ISSN: 0026-9247

DOCUMENT TYPE: Journal LANGUAGE: German

OTHER SOURCE(S): CASREACT 104:88492

GRAPHIC IMAGE:

$$\begin{array}{c|cccc}
R & NH_2 & & H & R^1 \\
N & & & & N & R^1 \\
N & & & & & N
\end{array}$$

ABSTRACT:

The reactions of RNHC(:NH)NH2 (R = Me, hexyl, cyclohexyl, CH2Ph, CH2CH2Ph, Ph, 1-naphthyl) with Me acrylate in DMF or EtOH preferentially afford dihydropyrimidinones I. The structures of I (R = hexyl, CH2Ph, Ph) were proved by comparison with authentic samples, which were prepared from RNHCH2CH2CO2Et and H2NCN. I (R = Ph) is not identical with authentic 2-phenylamino-5,6-dihydro-4-pyrimidinone (prepared from 2-methylthio-5,6-dihydro-4-pyrimidinone). HN:CR1NH (R1 = pyrrolidino, hexamethyleneimino, morpholino, N-methylpiperazino) react with Me acrylate in DMF to afford dihydropyrimidinones II. Action of morpholine-4-carboxamidine on Me acrylate in EtOH yields II (R1 = morpholino) as byproduct and EtOCH2CH2CON:CR1NH2 as main product.

=> DIS L2 7 IBIB IABS

THE ESTIMATED COST FOR THIS REQUEST IS 2.74 U.S. DOLLARS DO YOU WANT TO CONTINUE WITH THIS REQUEST? (Y)/N:Y

L2 ANSWER 7 OF 34 CAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 1985:24345 CAPLUS

DOCUMENT NUMBER: 102:24345

TITLE: Borane reduction of uracil derivatives

AUTHOR(S): Ghosh, Chandrakanta; Schmidt, Diane Grob; Pal, Bimal

С.

CORPORATE SOURCE: Inf. Div., Oak Ridge Natl. Lab., Oak Ridge, TN, 37831,

USA

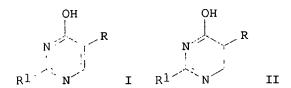
SOURCE: Journal of Organic Chemistry (1984), 49(26), 5256-7

CODEN: JOCEAH; ISSN: 0022-3263

DOCUMENT TYPE: Journal LANGUAGE: English

OTHER SOURCE(S): CASREACT 102:24345

GRAPHIC IMAGE:



ABSTRACT:

The reduction of uracil derivs. with BH3-THF was studied. Pyrimidinols I (R = CO2H, R1 = SH, OH) undergo reduction of the 5,6-double bond with concomitant decarboxylation to form dihydropyrimidines II (R = H, R1 = SH, OH) in 77 and 27% yield, resp. I (R = CO2Et, R1 = SH, SCH2Ph) form II (R = CO2Et, R1 = SH) and I (R = CH2OH, R1 = SCH2Ph) in 76 and 42 % yield, resp. Orotic acid and I (R = CO2H, R1 = SCH2Ph; R = H, R1 = SH; R = F, iodo, R1 = OH) did not react. The presence of an electron-withdrawing functional group such as CO2Et at C-5 of the pyrimidine ring facilitates reduction of the 5,6-double bond. I (R = CH2OH, R1 = SCH2Ph) was debenzylated to I (R = CH2OH, R1 = SH) with Na-NH3.

=> DIS L2 8 IBIB IABS THE ESTIMATED COST FOR THIS REQUEST IS 2.74 U.S. DOLLARS DO YOU WANT TO CONTINUE WITH THIS REQUEST? (Y) /N:Y

ANSWER 8 OF 34 CAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 1980:57902 CAPLUS

DOCUMENT NUMBER:

92:57902

TITLE:

Structure and tautomerism of the neutral and

monoanionic forms of 2-thiouracil, 2,4-dithiouracil,

their nucleosides, and some related derivatives

AUTHOR(S):

Psoda, Anna; Shugar, David

CORPORATE SOURCE:

Inst. Exp. Phys., Univ. Warsaw, Warsaw, 02-089, Pol.

SOURCE:

Acta Biochimica Polonica (1979), 26(1-2), 55-72

CODEN: ABPLAF; ISSN: 0001-527X

DOCUMENT TYPE:

Journal English

LANGUAGE: GRAPHIC IMAGE:

For diagram(s), see printed CA Issue.

ABSTRACT:

UV and IR data indicate that the monoanionic form of aqueous 2-thiouracil exists as the .apprx.1:1 I-II tautomeric equilibrium mixture; the charge is localized on the O atom in contrast to the charge delocalization in 2,4-dioxopyrimidines or 4-thiouracil monoanions. Neutral 2,4-dithiouracil and 2,4-dithiouridine exist in the dithione form in both aqueous and nonaq. media. The monoanion of 2,4-dithiouracil exists as the tautomer mixture 95:5 III-IV; the monoanion tautomers are delocalized as is the monoanion of dithiouridine. 2-(Methylthio)-4-pyrimidinone exists as V in CHCl3 and as a .apprx.1:1 V-VI equilibrium mixture in H2O.

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DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)	SINCE FILE ENTRY	TOTAL SESSION
CA SUBSCRIBER PRICE	-6.00	-6.00

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COST IN U.S. DOLLARS	SINCE FILE ENTRY	TOTAL SESSION
FULL ESTIMATED COST	0.06	36.76
DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)	SINCE FILE ENTRY	TOTAL SESSION
CA SUBSCRIBER PRICE	0.00	-6.00

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L2 ANSWER 9 OF 34 CAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 1979:104940 CAPLUS

DOCUMENT NUMBER: 90:104940

TITLE:

Mercaptoheterocyclic resin stabilizers

INVENTOR(S):

Sekiguchi, Tetsuo; Abe, Masami; Tsuruga, Kouji;

Tominaga, Nobuhide

PATENT ASSIGNEE(S):

Argus Chemical Corp., USA

SOURCE:

U.S., 12 pp. CODEN: USXXAM

DOCUMENT TYPE:

Patent

LANGUAGE:

English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.		DATE
				-	
US 4105627	Α	19780808	US 1976-725356		19760920
PRIORITY APPLN. INFO.:			US 1976-725356	Α	19760920
ABSTRACT:					

Mercaptan derivs. of heterocyclic amines in combination with epoxides, organic phosphites, or metal salts of fatty acids are heat stabilizers for polymers. Thus, poly(vinyl fluoride) [24981-14-4] 100, tris(nonylphenyl) phosphite 0.3, and 2-mercapto-4-hydroxy-6-methylpyrimidine (I) [56-04-2] 0.1 part were mixed on a 2-roll mill 3 min at 210° and molded 5 min at 250° and 100 kg/cm2. The sheet was colorless, compared with carbonized and brittle in the

kg/cm2. The sheet was colorless, compared with carbonized and brittle in the absence of I.

=> DIS L2 10 IBIB IABS

THE ESTIMATED COST FOR THIS REQUEST IS 2.74 U.S. DOLLARS DO YOU WANT TO CONTINUE WITH THIS REQUEST? (Y)/N:Y

L2 ANSWER 10 OF 34 CAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER:

1976:561303 CAPLUS

DOCUMENT NUMBER:

85:161303

TITLE:

Heat stabilizers for vinyl halide resins

INVENTOR(S):

Sekiguchi, Tetsuo; Abe, Masami; Tsuruga, Koji;

Tominaga, Nobuhide

PATENT ASSIGNEE(S):

Adeka Argus Chemical Co., Ltd., Japan

SOURCE:

Jpn. Kokai Tokkyo Koho, 14 pp.

CODEN: JKXXAF

DOCUMENT TYPE: LANGUAGE: Patent Japanese FAMILY ACC. NUM. COUNT: 1 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.		DATE
				-	
JP 51088542	A2	19760803	JP 1975-13797		19750131
PRIORITY APPLN. INFO.:			JP 1975-13797	Α	19750131
ABSTRACT:					

PVC [9002-86-2] (optionally containing ABS [9003-56-9]) and poly(vinyl fluoride) [24981-14-4] contained heterocyclic compds. containing -N:C(SH)NRC(:R1)- groups (R = H, alkyl, aryl; R1 = O, S; including tautomeric forms or salts) as heat stabilizers. For example, a PVC composition containing DOP 50, epoxidized soybean oil

2, tris(nonylphenyl) phosphite 0.5, stearic acid 0.5, and 3,5-dimercapto-1,2,4-triazole (I) [5650-03-3] 0.05 phr had heat stability (175°) 60 min, compared with 30 min for a control not containing I.

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	ENTRY	SESSION
FULL ESTIMATED COST	5.94	42.70
DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)	SINCE FILE	TOTAL
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CA SUBSCRIBER PRICE	-1.50	-7.50

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COST IN U.S. DOLLARS	SINCE FILE ENTRY	TOTAL SESSION
FULL ESTIMATED COST	0.06	42.76
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CA SUBSCRIBER PRICE	0.00	-7.50

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ANSWER 11 OF 34 CAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 1976:547731 CAPLUS

DOCUMENT NUMBER:

85:147731

TITLE:

Studies on the inhibition of metal corrosion. III. Some thiocarbonyl compounds as corrosion inhibitors

for copper in acid solution

AUTHOR(S):

CORPORATE SOURCE:

Hirooka, Motoko; Zen, Shonosuke Sch. Pharm. Sci., Kitasato Univ., Tokyo, Japan

SOURCE:

Nippon Kagaku Kaishi (1976), (7), 1167-70

DOCUMENT TYPE:

CODEN: NKAKB8; ISSN: 0369-4577

LANGUAGE:

Journal Japanese

ABSTRACT:

Thiocarbonyl compds. were effective corrosion inhibitors in acidic media. The most effective compound was 2-mercaptobenzimidazole [583-39-1] which had 92.1 and 100% inhibitive effect at the concentration of 32 and 250 ppm, resp. The effect was due to the formation of an insol. chelate film on the Cu surface. The chelate structure was identified by ir spectra.

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COST IN U.S. DOLLARS	SINCE FILE	TOTAL
	ENTRY	SESSION
FULL ESTIMATED COST	3.20	45.96
DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)	SINCE FILE ENTRY	TOTAL SESSION
CA SUBSCRIBER PRICE	-0.75	-8.25

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FILE CONTAINS CURRENT INFORMATION. LAST RELOADED: Apr 21, 2006 (20060421/UP).

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COST IN U.S. DOLLARS	SINCE FILE ENTRY	TOTAL SESSION
FULL ESTIMATED COST	0.06	46.02
DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)	SINCE FILE ENTRY	TOTAL SESSION
CA SUBSCRIBER PRICE	0.00	-8.25

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L2 ANSWER 12 OF 34 CAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 1976:432949 CAPLUS

DOCUMENT NUMBER: 85:32949

TITLE: Cyclization of N-(2-cyanoethyl)thioureas to

5,6-dihydro-2-thiouracils

AUTHOR(S): Dembecki, Marga; Pyl, Theodor

CORPORATE SOURCE: Sekt. Chem., Ernst-Moritz-Arndt-Univ., Greifswald,

Ger. Dem. Rep.

SOURCE: Zeitschrift fuer Chemie (1976), 16(4), 148-50

CODEN: ZECEAL; ISSN: 0044-2402

DOCUMENT TYPE: Journal

LANGUAGE: German

OTHER SOURCE(S): CASREACT 85:32949

GRAPHIC IMAGE:

ABSTRACT

Dihydrothiouracils I (R = H, Me, Et, CHMe2, Bu, Ph, CH2Ph; R1 = H, Me, Ph) (19 compds.) were prepared in 25-100% yields by cyclization of R1NHCSNRCH2CH2CN by heating with dilute or concentrated HCl. Refluxing I (R or R1 = H) with BzCl in pyridine gave 20-100% I (R = Me, Et, CHMe2, Bu, Ph, CH2Ph, R1 = Bz; R = Bz, R1 = Me, Ph).

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    ANSWER 1 OF 1 CAPLUS COPYRIGHT 2006 ACS on STN
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    2004:120553 CAPLUS
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    Bicyclic thioureas as lubricating oil antiwear, anticorrosion, and
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    antioxidant additives to replace zinc dialkyl dithiophosphates
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    Mukkamala, Ravindranath
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    U.S. Pat. Appl. Publ., 4 pp.
    CODEN: USXXCO
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